RECOMBINATION IN BACTERIA
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Recombination has been ascertained in a group of $E.\,coli$ strains, one of which, strain K-12, has been studied more extensively than any other. In this strain, recombination is governed by a system of mating types which shows an exceptional mode of hereditary transmission. Original K-12 is self-compatible (F+); some mutants derived from it proved to be self-incompatible (F-). Crosses $F+\times F-$ are fertile, and all the progeny is F+. An F- strain can be converted to F+ by incubation in certain conditions with an F- strain, no other genetic exchange taking place with the transfer of F+. The nature of the agent transducing self-compatibility is unknown. The kinetics of transduction seems to indicate that contact between cells is essential. An originally F- strain which has become F+ by transduction is stable and can transduce F+ in its turn.

Two other types of behaviour for compatibility have been found. A strain (Hfr) which behaves as an F+ for compatibility, shows much higher fertility (i.e., frequency of recombination) than comparable F+ strains; but cannot transduce F+. The progeny of an $Hfr\times F-$ cross is mostly F-, but a few recombinants are Hfr, in close linkage with a Gal marker. Another strain (Fr) behaves as an F-, but cannot be transduced to F+. Progeny from $F+\times Fr$ shows a segregation of F+ and F-. Fr strains do not compete with other F- strains for F+ when a limited supply of this agent is available.

The mating-type system shows a profound influence on the recombination pattern, as shown by the segregation of unselected markers. The contribution from the F + (or Hfr) parent undergoes an almost constant elimination, which is highest for one particular point or region of the chromosome (s). This results in considerable differences of reciprocal crosses.

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